Appl. No. 09/929,863 Amdt. Dated August 21, 2003 Reply to Office action of March 12, 2003

IN THE CLAIMS:

Claim 1. (Currently amended) A method to induce differentiation of a <u>an isolated or purified</u> naïve CD4⁺ T cell to a Tr1 cell comprising contacting the <u>naïve CD4⁺</u> T cell with an appropriate amount of interferon-α (IFN-α) <u>and an appropriate amount of IL-10</u>.

Claim 2. (Currently amended) The method of Claim 1, wherein said Tr1 cell is characterized by:

- a) CD4 expression;
- b) high levels of IL-10 production;
- c) significant levels of TGF-β or IFN-γ production; and
- d) little or no production of IL-4 or IL-2.

Claim 3. (Currently amended) The method of Claim 2, wherein:

- a) said high level of the IL-10 production is at least 6000 pg in 1 ml for 10⁶ colls in 48 h;
- b) said significant level of the TGF-β production is at least 600 100 pg in 1 ml for 10⁶ cells in 48 h;
- c) said significant level of the IFN-γ production is at least 1000 400 pg in 1 ml for 10⁶ cells in 48 h;
- d) said little or no the IL-4 production is less than 200 pg in 1 ml for 10⁶ sell in 48 h; or
- e) said little or no the IL-2 production is less than 200 pg in 1 ml for 10⁶ cell in 48-h;

when evaluated from cultures of about 106 cells per ml per 48 hours.

Appl. No. 09/929,863 Amdt. Dated August 21, 2003 Reply to Office action of March 12, 2003

- Claim 4. (Currently amended) The method of Claim 2, wherein:
 - a) said high level-of the IL-10 production is at least 6000 12000 pg in 1 ml for 106-cells in 48 h:
 - b) said significant level of the TGF-β production is at least 600 pg in 1 ml for 10⁶ cells in 48-h;
 - c) said significant level of the IFN-γ production is at least 1000 pg in 1 ml for 10⁶ cells in 48-h;
 - d) said little or no the IL-4 production is less than 200 100 pg in 1 ml for 10⁶ cell in 48-h; or
 - e) said little or no the IL-2 production is less than 200 100 pg in 1 ml for 10⁶ cell-in 48 h;

when evaluated from cultures of about 10⁸ cells per ml per 48 hours.

- Claim 5. (Original) The method of Claim 2, wherein said Tr1 cell:
 - a) has a reduced proliferative potential in response to polyclonal activation;
 and/or
 - b) suppresses response to alloantigens by responder T cells.
- Claim 6. (Currently amended) The method of Claim 1, wherein said Tr1 cells <u>cell</u> suppress<u>es</u> antigen-specific activation of <u>a</u> naive autologous T cells <u>cell</u>.
- Claim 7. (Original) The method of Claim 5, wherein said suppressed response to alloantigens is mediated by IL-10 and/or TGF-β.

Claims 8-10. (Cancelled).

Claim 1/1. (Original) The method of Claim 1, wherein said contacting is in combination with an antigen.

Claim 1/2. (Original) The method of Claim 1/1, wherein said antigen is an alloantigen.

Atty. Docket No. DX01135

Appl. No. 09/929,863 Amdt. Dated August 21, 2003 Reply to Office action of March 12, 2003

10

Claim 1/3. (Currently amended) The method of Claim 1, wherein said Tr1 cells are cell is further proliferated in IL-15.

11

Claim 14. (Currently amended) The method of Claim 1, wherein said Tr1 cells are cell is further tested for antigen specificity.

Claims 15-18. (Cancelled).

Claim 19. (New) A method to induce differentiation of an isolated or purified cord blood T cell to a Tr1 cell comprising contacting the cord blood cell with an appropriate amount of IFN-α.